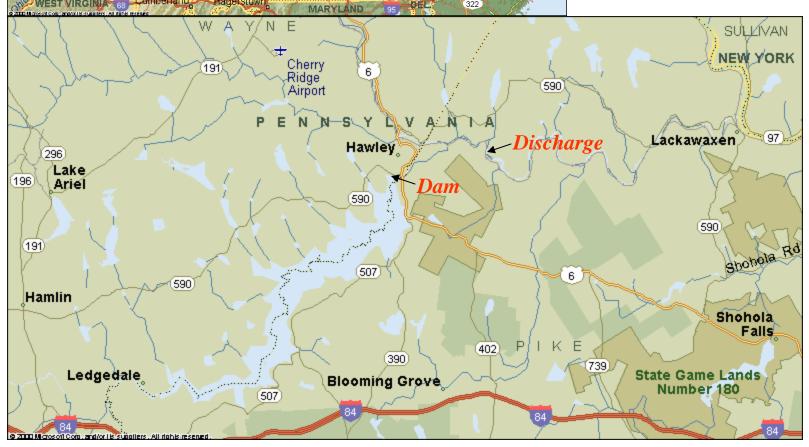


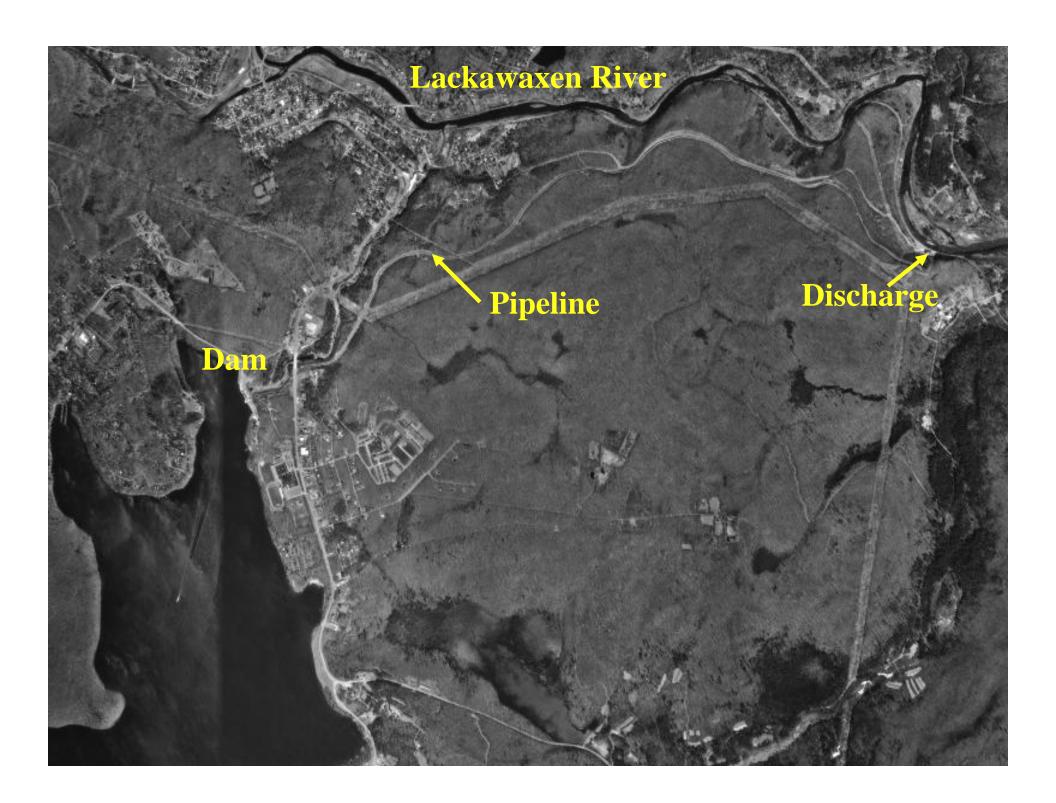


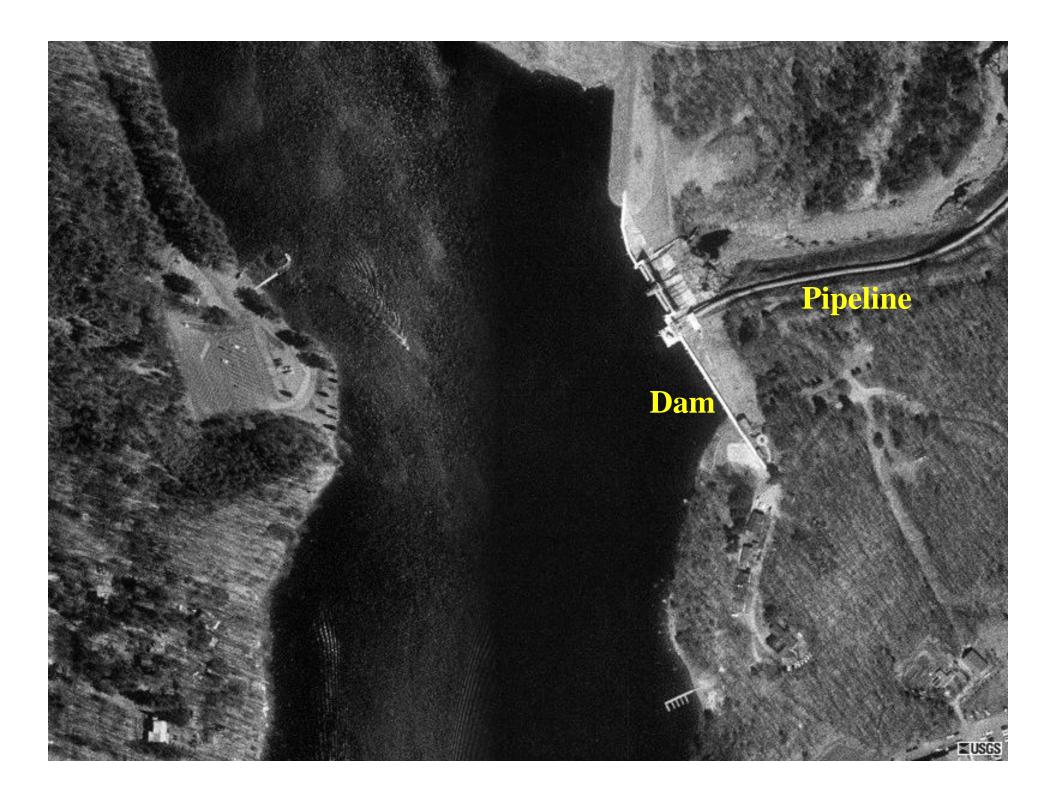
Lake Wallenpaupack:

- 5,700 acres
- 116,650 acre-feet*
- 40 ft mean depth
- pipeline 3.5 mi long

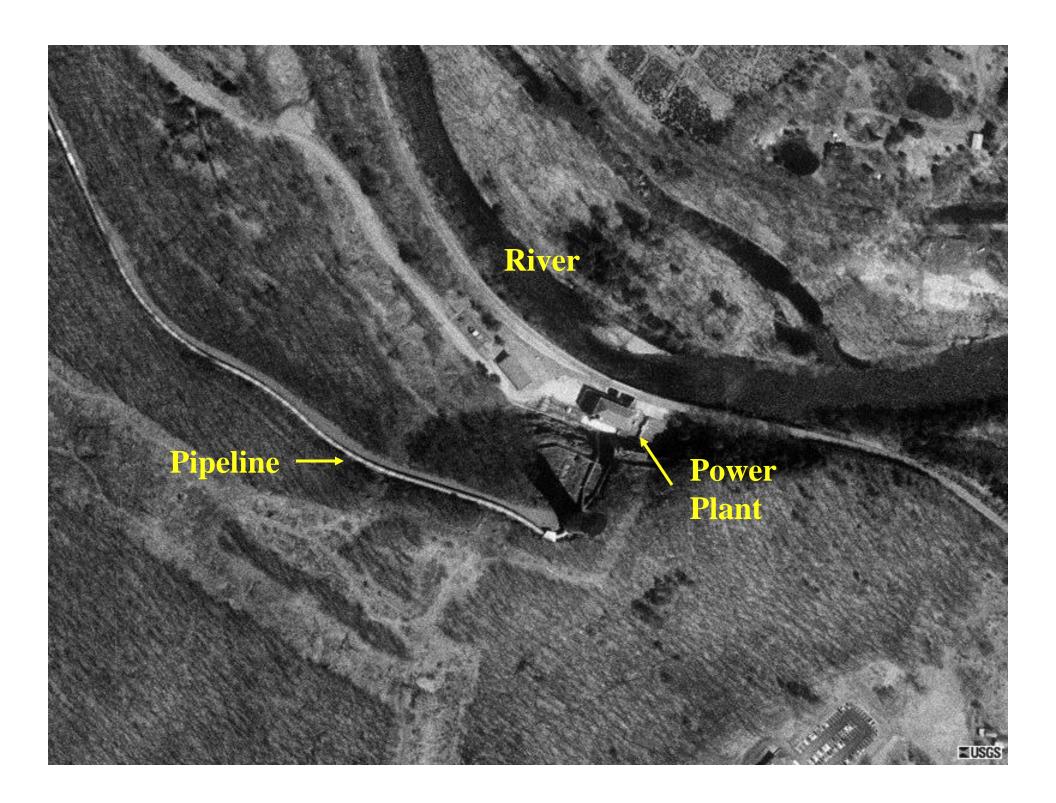


* usable storage

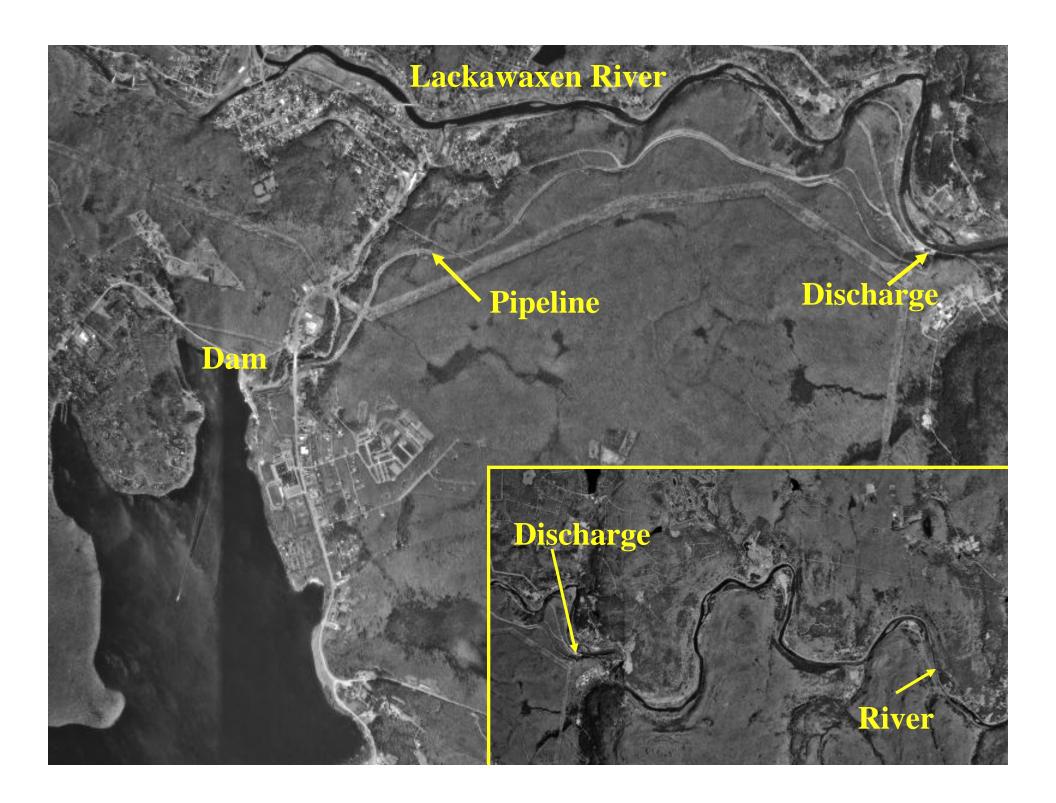




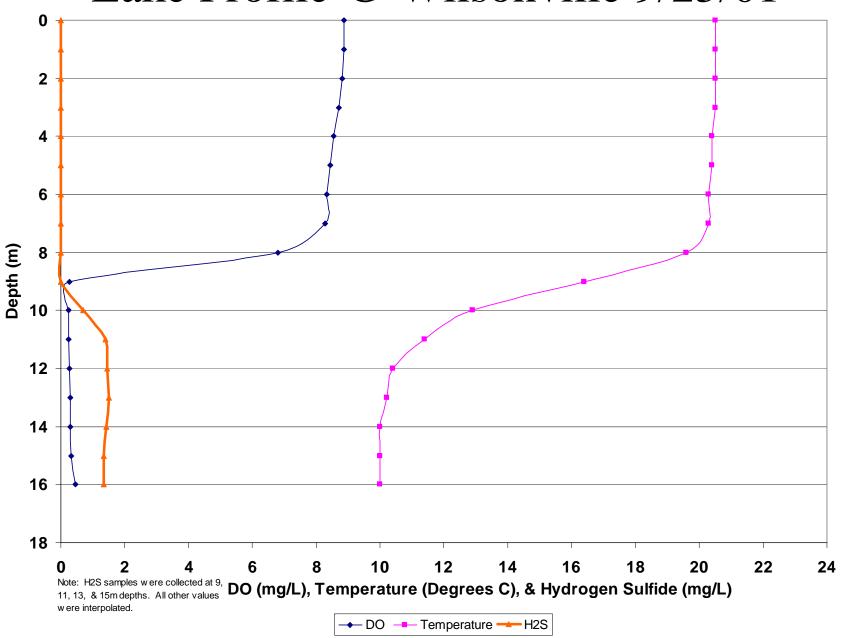








Lake Profile @ Wilsonville 9/25/01



Nature of the Problem

- Sulfate sources: precipitation, inflow, septics
- <u>Hypolimnetic anoxia</u> during summer enhanced by organic matter loading
- Sulfate reduced to <u>hydrogen sulfide</u> during anoxia
- <u>Hypolimnetic withdrawal</u> during operation (supports coldwater fishery) results in tailrace degassing of H₂S
- Need to either <u>prevent formation or enhance</u> removal of H₂S to minimize odor problem

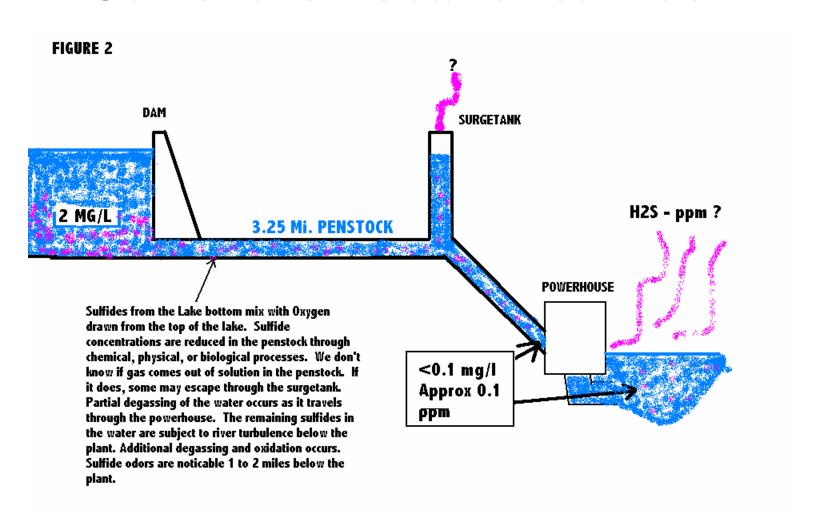
Expert Panel Gathering

- Robert Wetzel, Univ. North Carolina
- Robert Gambrell, Louisiana State
- Steve Ashby, COE Vicksburg
- George Luther, Univ. Delaware
- Richard Ruane, Reservoir Env. Mgmt., Inc.
- Forest Dierberg, DB Environmental, Inc.
- Yuefeng Xie, Penn State Harrisburg
- Frank Browne, FX Browne, Inc.

Objectives

- Short-Term:
 - In-lake or in-pipeline H₂S control
 - prevent formation
 - enhance oxidation or precipitation
- Long-Term:
 - Watershed management
 - reduce sulfate inputs
 - reduce organic matter loading

Current removal of sulfide





In-Pipe Approaches

- Enhance mixing
 - Immediate dilution effects
 - Longer time-of-travel for oxidation by DO
- Promote oxidation by dissolved oxygen
 - Air bubbler at pipeline intake
 - Add oxygen to air bubbler
- pH modification, chemical oxidation
 - Increase pH (form HS⁻, faster oxidation
 - Add H₂O₂ or Cl⁻ to oxidize H₂S

H₂S Oxidation by Dissolved Oxygen

May be too slow for the pipeline

Time (min)	Sulfide (mg/L)		
0	0.23		
1	0.23		
3	0.23		
5	0.23		
10	0.22		
30	0.20		

DO = 7.6

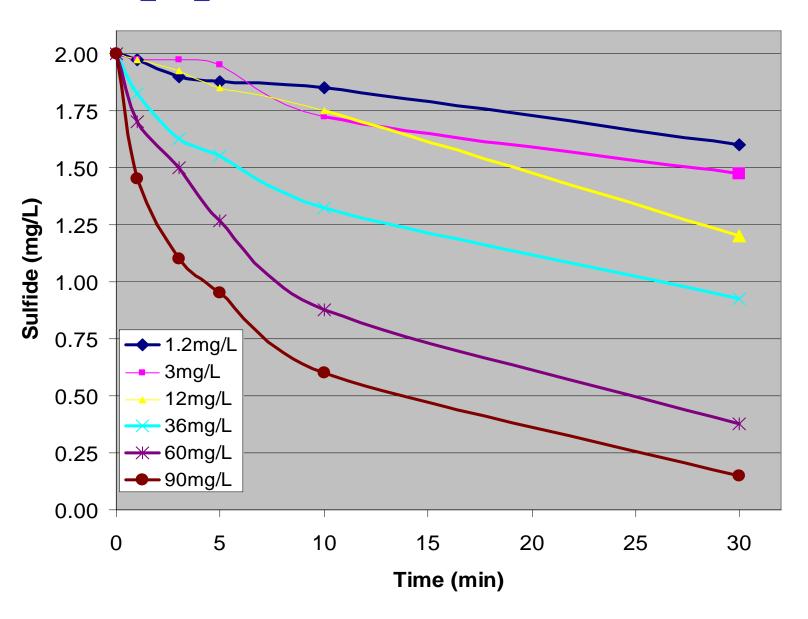
pH Modification Chemical Oxidation

HS⁻ predominant at pH > 7.0 (no odor) S_8 produced at pH < 7 with oxidation

 $4x H_2 O_2$ needed at pH > 7.5 2:1 mole ratio $H_2 O_2$ recommended at pH = 7 15-minute $H_2 O_2$ residence time suggested

Oxidants H₂0₂ and NaOCl tested

H₂O₂ as Oxidant with Lake Water



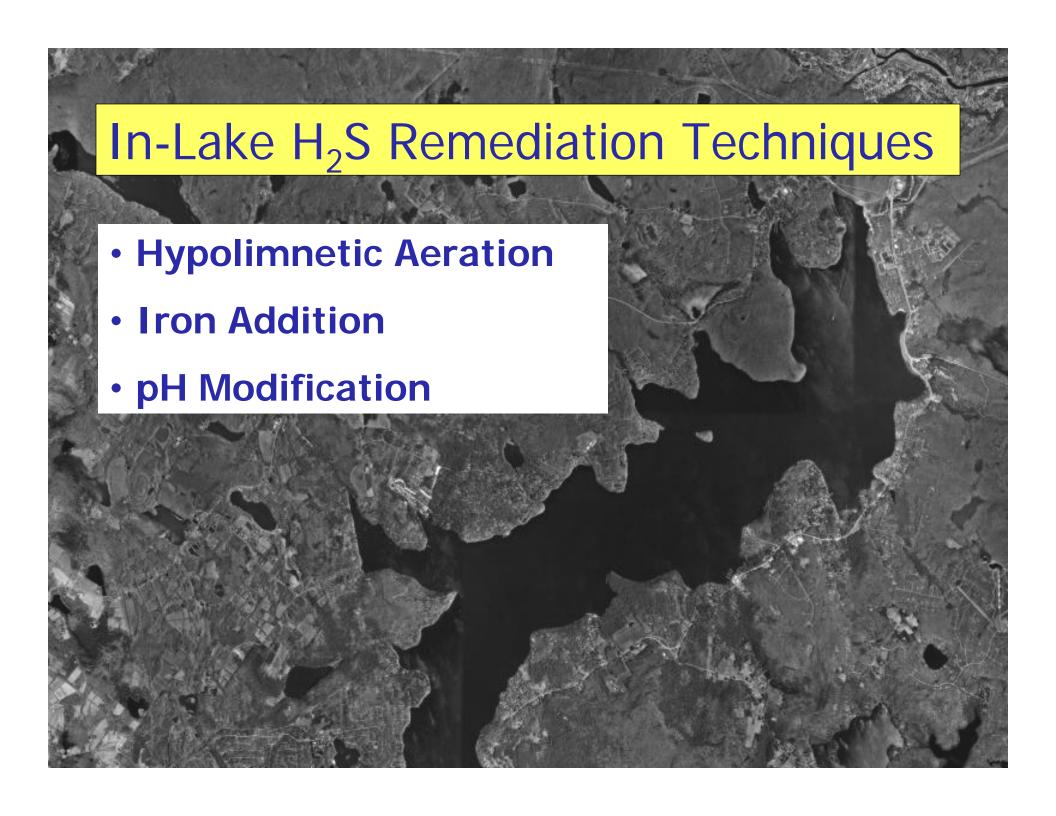
NaOCI as Oxidant with Lake Water

30-minute exposure

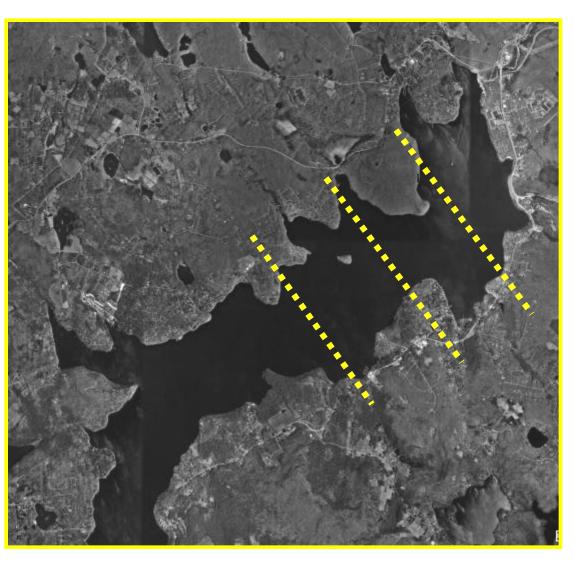
Concentrations in mg/L

pH = 6.58

NaOCl	Initial H ₂ S	Final H ₂ S	Free Cl
1	1.725	0.875	$oldsymbol{0}$
$\frac{1}{2}$	1.723	0.875	0 - 0
5	1.825	0.625	0



Hypolimnetic Aeration

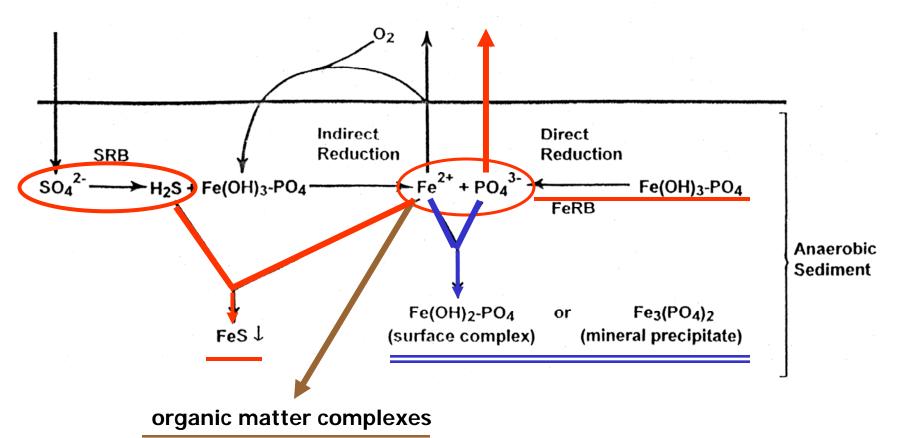


- Maintain thermocline (cold water)
- Enhance oxidation of H₂S by oxygen
- Lake volume to treat (oxid. rates)
- Avoid nutrient upwelling

In-Lake Iron Addition

- Under oxic conditions:
 - -Fe²⁺ oxidized to Fe³⁺
 - $-Fe^{3+}$ can oxidize H_2S to S_8 at pH > 6
- Under anoxic conditions:
 - -Fe³⁺ is reduced to Fe²⁺
 - -Fe²⁺ and H₂S can form FeS_{aq} and FeS₂↓
 - -PO4³- can be released with insufficient Fe

Sulfur - Iron- Phosphorus Cycle Interactions



Modified from: Wetzel, 2001

Modified from: Roden and Edmonds, 1997

In-Lake pH Modification

- Add limerock in area near intake:
 - to increase pH to about 7.5
 - favors HS⁻ over H₂S at higher pH's
 - $-HS^-$ predominant at pH > 7.0 (no odor)
- Treatment area and longevity unknown

Watershed Management Efforts

- Reduce sulfate/sulfur input
- Reduce organic matter loading
- Reduce nutrient loading
- Long-term solution

Hawley

Lackawaxen River Watershed

Summary of Techniques

- In-Pipe Remediation:
 - Enhance mixing
 - Promote oxidation by oxygen
 - pH modification
 - Chemical oxidation (H₂O₂ or NaOCl)
- In-Lake Remediation:
 - Hypolimnetic aeration
 - Iron addition
 - pH modification
- Watershed Remediation
 - Nutrient and organic matter reductions
 - Sulfate/sulfur loading reductions